

**UNITED STATES DEPARTMENT OF COMMERCE****Patent and Trademark Office**Address: COMMISSIONER OF PATENTS AND TRADEMARKS  
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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. |
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08/976, 566 11/24/97 POTTER

A 9001-0016.01

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| ROBINS & ASSOCIATES<br>90 MIDDLEFIELD ROAD<br>SUITE 200<br>MENLO PARK CA 94025 | HM22/0607 | EXAMINER |
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HM22/0607

EXAMINER

| GRASER, T | ART UNIT | PAPER NUMBER |
|-----------|----------|--------------|
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1641  
DATE MAILED:

06/07/00

19

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

|                              |                                      |                                      |
|------------------------------|--------------------------------------|--------------------------------------|
| <b>Office Action Summary</b> | Application No.<br><b>08/976,566</b> | Applicant(s)<br><b>Potter et al.</b> |
|                              | Examiner<br><b>Graser, Jennifer</b>  | Group Art Unit<br><b>1641</b>        |

Responsive to communication(s) filed on Request for CPA, 5/8/00

This action is **FINAL**.

Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11; 453 O.G. 213.

A shortened statutory period for response to this action is set to expire 3 month(s), or thirty days, whichever is longer, from the mailing date of this communication. Failure to respond within the period for response will cause the application to become abandoned. (35 U.S.C. § 133). Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

#### Disposition of Claims

Claim(s) 37, 40, 41, 44, and 45 is/are pending in the application.

Of the above, claim(s) \_\_\_\_\_ is/are withdrawn from consideration.

Claim(s) \_\_\_\_\_ is/are allowed.

Claim(s) 37, 40, 41, 44, and 45 is/are rejected.

Claim(s) \_\_\_\_\_ is/are objected to.

Claims \_\_\_\_\_ are subject to restriction or election requirement.

#### Application Papers

See the attached Notice of Draftsperson's Patent Drawing Review, PTO-948.

The drawing(s) filed on \_\_\_\_\_ is/are objected to by the Examiner.

The proposed drawing correction, filed on \_\_\_\_\_ is  approved  disapproved.

The specification is objected to by the Examiner.

The oath or declaration is objected to by the Examiner.

#### Priority under 35 U.S.C. § 119

Acknowledgement is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d).

All  Some\*  None of the CERTIFIED copies of the priority documents have been

received.

received in Application No. (Series Code/Serial Number) \_\_\_\_\_.

received in this national stage application from the International Bureau (PCT Rule 17.2(a)).

\*Certified copies not received: \_\_\_\_\_

Acknowledgement is made of a claim for domestic priority under 35 U.S.C. § 119(e).

#### Attachment(s)

Notice of References Cited, PTO-892

Information Disclosure Statement(s), PTO-1449, Paper No(s). \_\_\_\_\_

Interview Summary, PTO-413

Notice of Draftsperson's Patent Drawing Review, PTO-948

Notice of Informal Patent Application, PTO-152

--- SEE OFFICE ACTION ON THE FOLLOWING PAGES ---

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## **DETAILED ACTION**

### ***Continued Prosecution Application***

1. The request filed on May 8, 2000 for a Continued Prosecution Application (CPA) under 37 CFR 1.53(d) based on parent Application No. 08/976,566 is acceptable and a CPA has been established. An action on the CPA follows.

### ***Double Patenting rejections***

2. A Terminal Disclaimer was received with the After Final Amendment filed January 12, 2000. Accordingly, the former Double Patenting rejections have been dropped.

### ***Claim Rejections - 35 USC § 103***

3. The following is a quotation of 35 U.S.C. § 103 which forms the basis for all obviousness rejections set forth in this Office action:

A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Subject matter developed by another person, which qualifies as prior art only under subsection (f) or (g) of section 102 of this title, shall not preclude patentability under this section where the subject matter and the claimed invention were, at the time the invention was made, owned by the same person or subject to an obligation of assignment to the same person.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. § 103, the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 C.F.R. § 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later

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invention was made in order for the examiner to consider the applicability of potential 35 U.S.C. § 102(f) or (g) prior art under 35 U.S.C. § 103.

4. Claims 37, 40, 41 and 45 are rejected under 35 U.S.C. 103(a) as being unpatentable over as being unpatentable over Potter (5,476,657) in view of Bell et al (5,114,711).

Potter discloses proteins and subunit antigens from *P. haemolytica* for use in stimulating immunity against respiratory diseases such as pneumonia, including shipping fever pneumoniae (see abstract). Vaccines comprising an immunogenic amino acid sequence of *P. haemolytica* leukotoxin, or an amino acid sequence substantially homologous and functionally equivalent thereto, and a pharmaceutically acceptable carrier are disclosed (column 3, lines 3-24). Potter discloses production of recombinant *P. haemolytica* leukotoxin and specifically recites the production of leukotoxin 352 or "LKT 352" (column 17, lines 54-58). It is disclosed that vaccination with LKT 352 in combination with a *P. haemolytica* saline extract significantly reduced bovine respiratory disease morbidity and bovine respiratory disease mortality as compared to treatment with a placebo (column 25, lines 43-48). It is also specifically disclosed that prior to immunization, it may be desirable to increase the immunogenicity of the particular *Pasteurella* protein, or an analog of the protein, by linking the antigenic peptide to a carrier (column 13, lines 10-15). It is disclosed that suitable carriers may be proteins, polysaccharides, VP6 polypeptides of rotaviruses, viral proteins (column 13, lines 9-50). However, Potter does not particularly exemplify chimeric proteins comprising a leukotoxin derived from *P. haemolytica* and a peptide hormone, such as gonadotropin releasing hormone (GnRH) or an epitope thereof.

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Bell et al. disclose the recombinant production of chimeric proteins which are composed of two covalently linked cell modulators in a linear polypeptide sequence. It is taught that the cell modulators are interferons, lymphokines or cytokines (abstract). Column 3, lines 60-64, define cell modulators as including lymphokines, monokines, peptide hormones, or peptide growth factors.

At the time the invention was made it was well known in the art to those of ordinary skill that many different protein combinations could be prepared recombinantly that produce chimeric proteins and it was also well established that various immune modulators have been conjugated to such things as antibodies, ligands, or hormones, to act as site-specific delivery agents (some of which also display functional activity of the cytotoxin). Potter specifically discloses that the immunogenicity of the *P. haemolytica* leukotoxin (a cytotoxin), and fragments thereof, could be made more immunogenic by linking it to a carrier such as proteins, polysaccharides, inactive virus particles and other large, slowly metabolized molecules (column 13). Bell et al. specifically disclose that cytotoxins and peptide hormones may be linked together to treat disease. It would have been obvious to one of ordinary skill in the art at the time the invention was made that a peptide hormone as disclosed by Bell et al. could be linked to at least one epitope of a leukotoxin derived from *P. haemolytica*, as taught by Potter, because the leukotoxin is a cytotoxin which Bell specifically teaches may be linked to peptide hormones, for a dual immune modulating effect. One of ordinary skill in the art would expect to increase the immune response to the leukotoxin and produce a more efficient vaccine against respiratory disease in ruminants by linking to an

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immune modulator. Potter specifically discloses that truncated leukotoxin, LKT 352, which lacks cytotoxic activity could be used as the *P.haemolytica* leukotoxin. Bell discloses the use of adjuvants and carriers and it would have been obvious to one of ordinary skill in the art to link the cytotoxin-peptide hormone to a carrier as the use of linking carriers to chimeric proteins was well known in the art for a means of increasing an immune response to an antigen. Although the use of gonadotropin releasing hormone is not specifically recited by Bell et al., it would have been an obvious choice in the cytotoxin-immunomodulator conjugates because it is a well known hormone which falls under the definition of "peptide hormone". These compositions would be structurally identical to those instantly claimed, i.e., a chimeric protein comprising leukotoxin coupled to a peptide hormone which is not a cytokine.

*Response to Applicants' arguments:*

Applicants have argued that the amendment to claim 37 which states that the leukotoxin polypeptide is coupled to a "selected peptide hormone which is not a cytokine" is sufficient to overcome the rejection. This argument has been fully and carefully considered but is not deemed persuasive because Bell et al discloses the use of immune-modulators other than cytokines in the cytotoxin conjugates. Column 3, lines 60-64, of Bell defines cell modulators as including lymphokines, monokines, peptide hormones, or peptide growth factors.

5. Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Group 1600 via the PTO Fax Center located in Crystal Mall 1. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The Group 1641 Fax number is (703) 308-4242 which is able to receive transmissions 24 hours/day, 7 days/week.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jennifer E. Graser whose telephone number is (703) 308-1742. The examiner can normally be reached on Monday-Friday from 7:00 AM-4:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, James Housel, can be reached on (703) 308-4027..

*Jennifer Graser*  
JENNIFER GRASER 6/4/02  
PATENT EXAMINER